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Entorhinal Cortex: Antemortem Cortical Thickness and Postmortem Neurofibrillary Tangles and Amyloid Pathology

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ABSTRACT

BACKGROUND AND PURPOSE: The entorhinal cortex, a critical gateway between the neocortex and hippocampus, is one of the earliest regions affected by Alzheimer disease–associated neurofibrillary tangle pathology. Although our prior work has automatically delineated an MR imaging–based measure of the entorhinal cortex, whether antemortem entorhinal cortex thickness is associated with postmortem tangle burden within the entorhinal cortex is still unknown. Our objective was to evaluate the relationship between antemortem MRI measures of entorhinal cortex thickness and postmortem neuropathological measures.

MATERIALS AND METHODS: We evaluated 50 participants from the Rush Memory and Aging Project with antemortem structural T1-weighted MR imaging and postmortem neuropathologic assessments. Here, we focused on thickness within the entorhinal cortex as anatomically defined by our previously developed MR imaging parcellation system (Desikan-Killiany Atlas in FreeSurfer). Using linear regression, we evaluated the association between entorhinal cortex thickness and tangles and amyloid- β load within the entorhinal cortex and medial temporal and neocortical regions.

RESULTS: We found a significant relationship between antemortem entorhinal cortex thickness and entorhinal cortex ($P = .006$) and medial temporal lobe tangles ($P = .002$); we found no relationship between entorhinal cortex thickness and entorhinal cortex ($P = .09$) and medial temporal lobe amyloid- β ($P = .09$). We also found a significant association between entorhinal cortex thickness and cortical tangles ($P = .003$) and amyloid- β ($P = .01$). We found no relationship between parahippocampal gyrus thickness and entorhinal cortex ($P = .31$) and medial temporal lobe tangles ($P = .051$).

CONCLUSIONS: Our findings indicate that entorhinal cortex–associated in vivo cortical thinning may represent a marker of postmortem medial temporal and neocortical Alzheimer disease pathology.

ABBREVIATIONS: AD = Alzheimer disease; EC = entorhinal cortex; SE = standard error

The human entorhinal cortex (EC) plays an integral role in memory formation and serves as the critical gateway between

the hippocampus and neocortex.¹ Located in the medial temporal lobe, the EC constitutes the anterior portion of the parahip-

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pocampal gyrus and is localized in vivo laterally by the rhinal sulcus, anteriorly by the amygdala and hippocampus, and posteriorly by the posterior portion of the parahippocampal gyrus.^{2,3} The EC is one of the earliest affected regions in Alzheimer disease (AD). Tau-associated neurofibrillary tangle pathology in AD follows a defined topographic and hierarchical pattern, first affecting the EC and then progressing to anatomically connected limbic and association cortices; in contrast, amyloid- β -associated pathology does not involve the EC in the earliest stages of AD but selectively affects the neocortical regions.^{4,5} Most important, the association of neuronal volume loss in the EC has been shown to parallel τ -associated tangle pathology (neurofibrillary tangles and neuritic plaques) but not senile plaques seen with amyloid- β deposition.⁴

Structural MR imaging provides visualization and quantification of volume loss and has been extensively investigated in AD.⁶⁻⁹ Using manually delineated assessments, early studies have shown that volumetric measures of the EC can identify individuals without dementia in the earliest stages of the AD process.⁷⁻⁹ Within the past decade, rapid advances in MR imaging postprocessing have led to the development of software tools for automatic quantification of human subcortical and neocortical regions.¹⁰ We have previously developed an MR imaging-based parcellation atlas for the human cerebral cortex, which has automatically delineated the entorhinal cortex.¹¹ The EC ROI from our parcellation atlas correlates with CSF levels of τ , amyloid,¹² and *Apolipoprotein E*¹³ and has been used to identify cognitively healthy¹⁴ and cognitively impaired individuals without dementia who are most likely to progress to clinical AD.^{15,16} However, whether our antemortem MR imaging-based measure of the EC is associated with established postmortem measures of AD pathology is still unknown.

In this study, we evaluated the relationship between antemortem MR imaging-based automated measurements of EC thickness and postmortem measures of neurofibrillary tangle and amyloid- β pathology. To assess the specificity of our EC ROI (anterior portion of the parahippocampal gyrus), we also evaluated the relationship between the thickness of the posterior parahippocampal gyrus and EC amyloid- β and τ pathology.

MATERIALS AND METHODS

Participants

We evaluated participants from the Rush Memory and Aging Project, a community-based longitudinal study of aging, which began in 1997.¹⁷ Details of the clinical and neuropathologic evaluations have been reported previously.^{17,18} Briefly, all participants underwent a uniform structured clinical evaluation that included a medical history, physical examination with emphasis on neurologic function, and neuropsychological testing (including the Mini-Mental State Examination and 20 other tests). All participants were evaluated in person by a neuropsychologist and a physician with expertise in the evaluation of older individuals with cognitive impairment. On the basis of physician evaluation and review of the cognitive testing and the neuropsychologist's opinion, participants were classified with respect to AD and other common conditions with the

Demographic information on participants in the current study^a

	Healthy (n = 25)	MCI (n = 18)	AD (n = 7)
% Female	64%	55%	71%
Education (yr)	15.6 (3.6)	14.9 (1.2)	13.2 (1.2)
Age at MRI (yr)	87.4 (5.0)	88.3 (5.4)	85.2 (4.4)
Age at death (yr)	90.3 (4.7)	91.1 (5.4)	88.7 (4.9)
Years between MRI and death	2.8 (1.3)	2.8 (1.2)	3.6 (1.5)
Entorhinal cortex thickness	1.50 (0.30)	1.49 (0.27)	1.40 (0.41)
EC tangle density	8.9 (6.2)	12.8 (10.3)	20.1 (12.9)
EC amyloid- β load	5.5 (5.1)	7.1 (5.1)	7.5 (4.9)

Note.—MCI indicates mild cognitive impairment.

^a All values are expressed as mean (SD).

potential to impact cognitive function according to the recommendations of the joint working group of the National Institute of Neurologic and Communicative Disorders and Stroke and the Alzheimer's Disease and Related Disorders Association.¹⁹ In this article, we focused on 50 participants, clinically defined at baseline (on study entry) as being cognitively healthy ($n = 25$), having mild cognitive impairment ($n = 18$), and having probable AD ($n = 7$) (Table), with concurrent antemortem MR imaging and postmortem neuropathologic assessments. The Rush University Medical Center institutional review board approved the study, and all participants gave written informed consent and signed an Anatomic Gift.

Imaging Assessments

We assessed previously obtained T1-weighted anatomic data by using a 1.5T MR imaging scanner (GE Healthcare, Milwaukee, Wisconsin). For the current study, all antemortem MR imaging data were acquired by using a 3D magnetization-prepared rapid acquisition of gradient echo sequence with the following parameters: TE = 2.8 ms, TR = 6.3 ms, preparation time = 1000 ms, flip angle = 8°, FOV = 24 × 24 cm, 160 sections, 1-mm section thickness, a 224 × 192 acquisition matrix reconstructed to 256 × 256, and 2 repetitions. The MR imaging data were automatically segmented with FreeSurfer 5.0 (<http://surfer.nmr.mgh.harvard.edu>; for additional details see McKhann et al¹⁹). Here, we focused on intracranial volume-corrected average thickness of the entire entorhinal cortex and posterior parahippocampal gyrus (average of the left and right hemispheres) as delineated with our previously developed automated cortical parcellation atlas (Desikan-Killiany atlas in FreeSurfer) (Fig 1).¹¹ In secondary analyses, we also evaluated baseline intracranial volume-corrected hippocampal volumes (average of the left and right hemispheres).²⁰

Neuropathologic Assessments

We used results from previously obtained neuropathologic evaluations and focused on amyloid and neurofibrillary tangle pathology within the entorhinal cortex and medial temporal (entorhinal and hippocampal) regions (for additional details on neuropathologic measures from the Rush Memory and Aging Project/Rush University Medical Center, please see Bennett et al^{21,22} and Barnes et al²³). Briefly, at least 2 tissue blocks from the entorhinal cortex and hippocampus (CA1/subiculum) were dissected from 1-cm coronal slabs fixed for 48–72 hours in 4% paraformaldehyde, embedded in paraffin, and cut into 20- μ m sections. Amyloid- β was labeled with MO0872 (1:100; Dako,

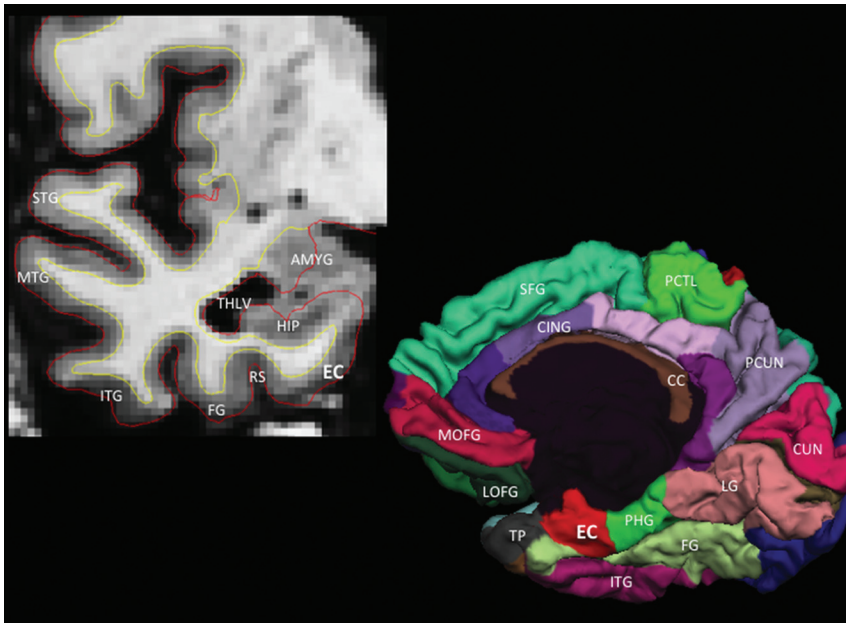


FIG 1. Coronal T1-weighted MR imaging illustrating the anatomic location of the entorhinal cortex, which is medial to the rhinal sulcus (RS) and fusiform gyrus (FG) and inferior to the hippocampus (HIP), temporal horn of the lateral ventricle (THLV), and amygdala (AMYG) (*upper left panel*). 3D cortical (pial) representation is of the right cortical (pial) surface, delineating the location of the entorhinal cortex on the medial hemisphere of the cerebral cortex (*lower right panel*). STG indicates superior temporal gyrus; MTG, middle temporal gyrus; ITG, inferior temporal gyrus; TP, temporal pole; PHG, parahippocampal gyrus; LG, lingual gyrus; CUN, cuneus cortex; PCUN, precuneus; FG, fusiform gyrus; PCTL, paracentral lobule; SFG, superior frontal gyrus; MOFG, medial orbitofrontal gyrus; LOFG, lateral orbitofrontal gyrus; CC, corpus callosum; CING, cingulate cortex.

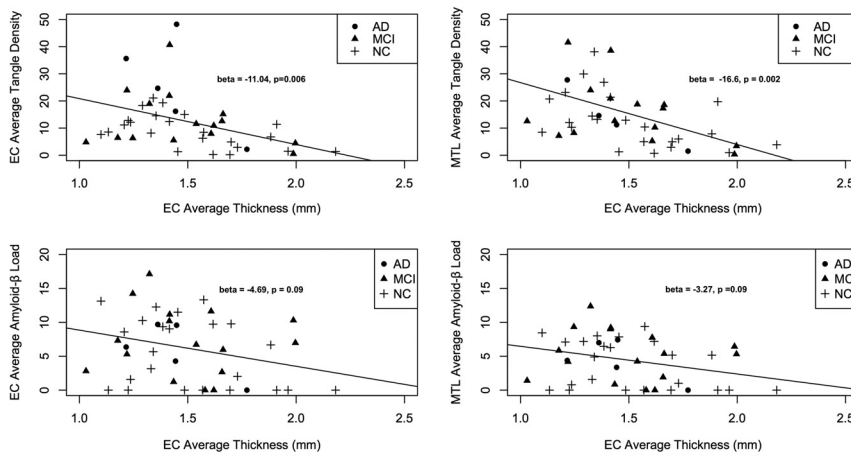


FIG 2. Scatterplots illustrating the relationship between average entorhinal cortex thickness and average tangle density within the entorhinal cortex (*upper left*) and medial temporal lobe (MTL, *upper right*), and average amyloid- β load within the EC (*lower left*) and medial temporal lobe (*lower right*). Best-fit regression line, β -coefficients, and P values from the logistic regression model are included (for additional details see the text).

Carpinteria, California), and paired helical filament τ was labeled with AT8 (1:800 in 4% horse serum; Innogenex, San Ramon, California), an antibody specific for phosphorylated τ . Images of amyloid- β -stained sections were captured for quantitative analysis by using a systematic random-sampling scheme, and calculation of the percentage area occupied by amyloid- β immunoreactive pixels was performed. Quantification of tangle density per square millimeter was performed with a stereologic mapping station. We used composite summary measures of the percentage area occu-

ried by amyloid- β and the density of neurofibrillary tangles by averaging the values for each lesion within the entorhinal cortex and medial temporal regions (entorhinal cortex and hippocampus) (for additional details see Bennett et al^{21,22} and Barnes et al²³). To evaluate neuropathology within the neocortex and to minimize multiple comparisons, we used a composite measure of tangle density and amyloid- β load within the midfrontal cortex, inferior temporal gyrus, inferior parietal cortex, calcarine cortex, cingulate region, and superior frontal gyrus (cortical tangle density and cortical amyloid- β load).

Statistical Analysis

Using linear regression, we evaluated the association between entorhinal cortex thickness and average tangle and amyloid- β load within the entorhinal cortex and medial temporal regions (entorhinal cortex + hippocampus). We also evaluated the relationship between entorhinal cortex thickness and cortical tangle density and amyloid- β load. In secondary analyses, we assessed the association between (posterior) parahippocampal gyrus thickness and tangles and amyloid- β load within the entorhinal cortex and medial temporal regions. In all analyses, we controlled for the effects of age at death, sex, and clinical diagnosis.

RESULTS

We found a relationship between ante-mortem entorhinal cortex thickness and postmortem tangle density within the entorhinal cortex (β -coefficient = -11.04 , standard error [SE] = 3.78 , P value = $.006$) and medial temporal regions (β -coefficient = -16.67 , SE = 5.04 , P = $.002$); lower EC thickness was associated with increased EC and medial temporal lobe tangle density (Fig 2). Even after controlling for the effects of hippocampal volume, the relationship between entorhinal cortex thickness and tangle density within the entorhinal cortex (β -coefficient = -9.36 , SE = 2.57 , P = $.03$) and medial temporal regions (β -coefficient = -12.89 , SE = 5.64 , P = $.02$) remained significant. We found no relationship between entorhinal cortex thickness and amyloid- β load within the entorhinal cortex (β -coefficient = -4.69 , SE = 2.70 , P = $.09$) and medial temporal regions (β -coefficient = -3.27 , SE = 1.90 , P = $.09$) (Fig 2).

We found a relationship between entorhinal cortex thickness and both cortical tangle density (β -coefficient = -37.5 , SE =

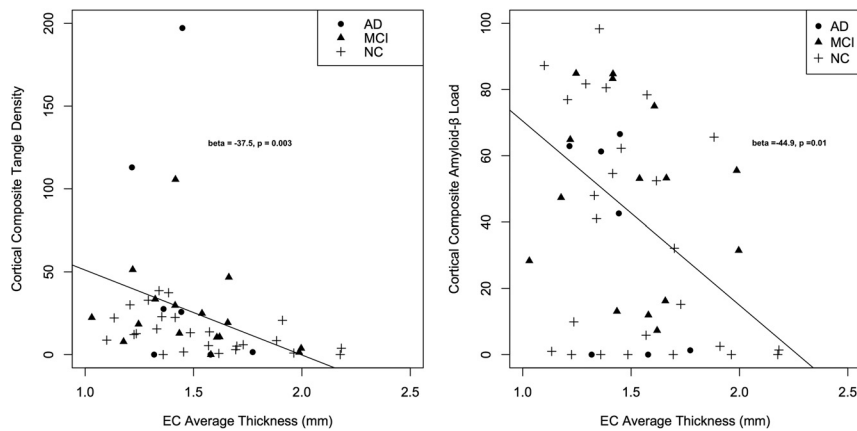


FIG 3. Scatterplots illustrating the relationship between average entorhinal cortex thickness and composite tangle density (*left*) and amyloid- β load (*right*) within the cerebral cortex (see text for details). Best-fit regression line, β -coefficients, and P values from the logistic regression model are included (for additional details see the text).

12.2, $P = .0003$) and amyloid- β load (β -coefficient = -44.9 , $SE = 17.2$, $P = .01$) (Fig 3). Even after we controlled for the effects of hippocampal volume, the relationship between entorhinal cortex and both cortical tangle density (β -coefficient = -34.4 , $SE = 7.9$, $P = .01$) and amyloid- β load (β -coefficient = -44.7 , $SE = 18.6$, $P = .02$) remained significant.

In contrast, we found no relationship between antemortem parahippocampal gyrus thickness and postmortem tangle density within the entorhinal cortex (β -coefficient = -4.04 , $SE = 4.33$, $P = .31$) and a trend toward significance for the medial temporal regions (β -coefficient = -11.52 , $SE = 5.71$, $P = .051$). Similarly, we found no relationship between parahippocampal gyrus thickness and amyloid load within the entorhinal cortex (β -coefficient = 1.02 , $SE = 2.71$, $P = .709$) and medial temporal regions (β -coefficient = 0.57 , $SE = 1.91$, $P = .766$). We also found no relationship between parahippocampal gyrus thickness and cortical tangle density (β -coefficient = -14.2 , $SE = 13.5$, $P = .30$) and amyloid- β load (β -coefficient = -8.7 , $SE = 18.6$, $P = .64$).

We performed subgroup analyses within our subset of cognitively healthy older participants ($n = 25$) to evaluate the relationship between MR imaging measures of entorhinal cortex thickness and neuropathology. Similar to our main results, we found a relationship between antemortem entorhinal cortex thickness and postmortem tangle density within the entorhinal cortex (β -coefficient = -10.32 , $SE = 4.03$, $P = .01$), medial temporal regions (β -coefficient = -14.48 , $SE = 6.61$, $P = .04$), and cortex (β -coefficient = -16.58 , $SE = 7.03$, $P = .02$). In contrast, we found no relationship between entorhinal cortex thickness and amyloid load either within the entorhinal cortex (β -coefficient = -5.04 , $SE = 3.91$, $P = .23$) or cortex (β -coefficient = -32.26 , $SE = 22.79$, $P = .12$).

DISCUSSION

Our results demonstrate that quantitative *in vivo* volumetric MR imaging measurements of the EC are associated with postmortem measures of entorhinal and neocortical AD pathology. Specifically, these results indicate that lower EC thickness predicts higher postmortem tangle load. We also found a similar association between EC thickness and postmortem tangle load

within the medial temporal regions (EC + hippocampus). Most important, this effect remained significant after controlling for hippocampal volume. Finally, we found a robust relationship between EC thickness and cortical tangle and amyloid- β pathology. Rather than representing a specific measure of EC tangle pathology, our combined findings suggest that an antemortem MR imaging measure of the entorhinal cortex likely captures Alzheimer-associated pathology within the medial temporal and neocortical regions.

Using neuropathologic assessments from the same individuals and building on our prior work,^{12,14} this study sug-

gests that EC-associated cortical thinning in AD may represent a marker of τ -associated pathology within both the entorhinal cortex and neocortex. Consistent with the known pattern of amyloid- β deposition in AD, which heavily involves the neocortex with relative sparing of the entorhinal cortex, an association was found between EC thickness and neocortical amyloid- β load but not entorhinal amyloid pathology, compatible with the hypothesis that amyloid deposition within the neocortex, rather than the EC may represent an early component of Alzheimer pathobiology.^{24,25} Our results also suggest that entorhinal cortex thickness may provide independent information about AD pathology even after accounting for hippocampal volume, further illustrating the importance of evaluating EC thickness as an early marker of *in vivo* Alzheimer neurodegeneration. Most interesting, even among cognitively healthy older adults, we found a selective relationship between entorhinal cortex thickness and tangle pathology but not amyloid pathology, suggesting the potential usefulness of quantitative MR imaging measures in preclinical AD.

Accumulating evidence suggests that τ pathology is closely associated with cognitive performance, particularly in the early stages of disease.^{26,27} These findings suggest that automated measures of entorhinal cortex atrophy may reflect regional τ pathology, which will be clinically useful for early AD detection and disease monitoring. Additionally, quantitative measures of EC and medial temporal structures may be combined with genetic, fluid (CSF or plasma), and cognitive parameters for risk stratification, which may become increasingly relevant for AD prevention and therapeutic trials. With the advent and use of novel agents for detecting *in vivo* τ deposition,²⁸ volumetric MR imaging could be integrated with PET imaging to determine whether regional measures of entorhinal atrophy and τ deposition provide independent or complementary information. Finally, beyond AD, automated assessments of the entorhinal cortex can be useful in other disorders of the medial temporal lobe such as medial temporal sclerosis, vascular dementia, and frontotemporal lobar degeneration.

A potential limitation of our study is that our imaging and neuropathology datasets were not coregistered; thus, this drawback limits the precise correspondence between antemortem and

postmortem definitions of the EC. Another limitation is the need for validation of our results in independent, community-based samples.

CONCLUSIONS

We found a strong association between an automated, antemortem MR imaging–based measure of EC thickness and postmortem neurofibrillary tangle burden within the entorhinal cortex, medial temporal lobe, and neocortex. We additionally detected a relationship between EC thickness and neocortical amyloid- β load. Considered together, our findings serve as a validation of our automated MR imaging measure of the EC and suggest that EC-associated cortical thinning in AD may represent a marker of medial temporal and neocortical AD neuropathology.

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